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FIG. 1A

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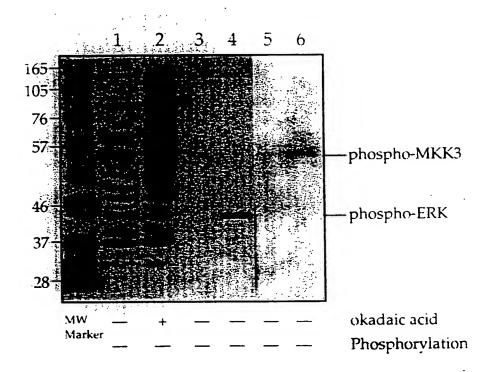
Production of Motif-Specific Context-Independent Antibodies Using Peptide Libraries as Antigent Comb, et al Atty. Docket No: CST-138 CIP2 Sheet 2 of 23

FIG. 1B

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30 phospho-Tyr peptides mixture	
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Production of Motif-Specific Context-Independent Antibodies Using Peptide Libraries as Antigens Comb, et al Atty. Docket No: CST-138 CIP2 Sheet 3 of 23

FIG. 1C



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FIG. 1D

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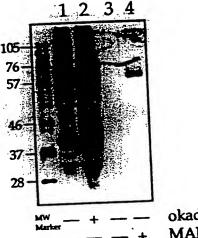
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Production of Motif-Specific Context-Independent Antibodies Using Peptide Libraries as Antig Comb, et al Atty. Docket No: CST-138 CIP2 Sheet 6 of 23

FIG. 2B



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okadaic acid MAPK phosphorylation

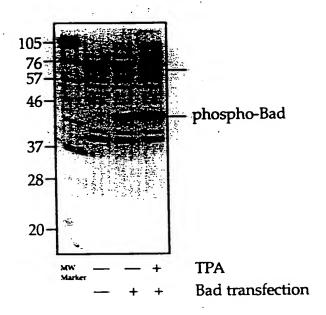
Production of Motif-Specific Context
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FIG. 3A

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Production of Motif-Specific Context-Independent Antibodies Using Peptide Libraries as Antigen Comb, et al Atty. Docket No: CST-138 CIP2 Sheet 8 of 23

FIG. 3B



Production of Motif-Specific Context
Production of Motif-Specific Context
Antibodies Using Peptide Libraries as Antigens

Comb, et al

Atty. Docket No: CST-138 CIP2

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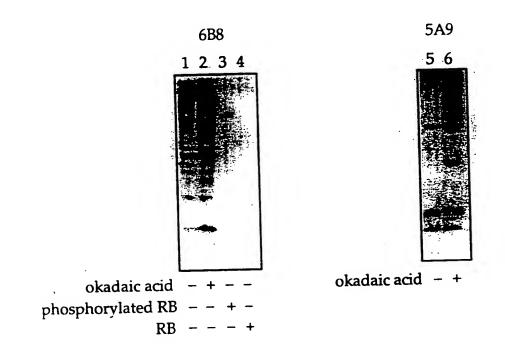
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FIG. 4A

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	CITION OF COURT			
ProXSer/ThrProXArg-P	XXXXXPro-X:8a-7mh-Pro-X-Arg/Lys-X-X-Cys	·	1.955	1.275
	(SEG ID NO.42)			
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Ser-P	**************************************		0.031	0.088
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	(SEQ ID NO.43)		٠.	
Rb (Thr373)-P	Val-lie-Pro-Pro-His-Thr-Pro-Val-Arg-Thr-Val-Met-Asn-Thr-Cys	hr-Cys	3.336	3.503
	(CEON CINCIPS)			
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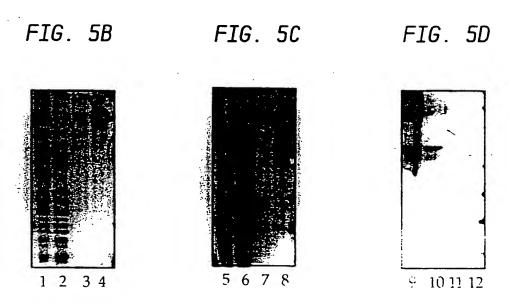
FIG. 4B



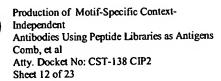
Production of Motif-Specific Context-Independent Antibodies Using Peptide Libraries as Antigen Comb, et al Atty. Docket No: CST-138 CIP2 Sheet 11 of 23

FIG. 5A

Acetylated BSA(ng) | BSA(ug) 0.5 2 10 50 200 500 10



Phospho-Akt Substrate Antibody



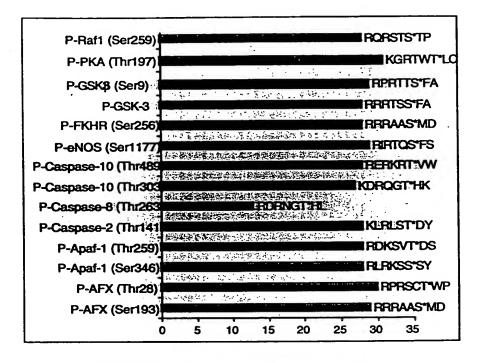
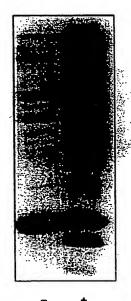


Figure 6: Signal to noise ratio of ELISA readings using Phospho-Akt Substrate Antibody with phosphopeptides of Akt substrates vs. nonphospho-peptides of Akt substrates.



calyculin A

Figure 7: Western analysis of calyculin A-treated A431 cells using Phospho-Akt Substrate Antibody.

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Production of Motif-Specific Context-Independent Antibodies Using Peptide Libraries as Antigens Comb, et al Atty. Docket No: CST-138 CIP2 Sheet 13 of 23

Phospho-PKA Substrate Antibody

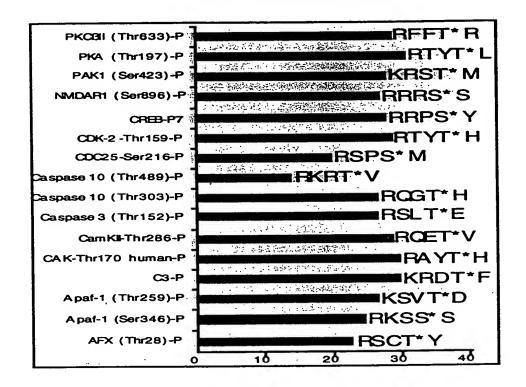


Figure 8: Signal to noise ratio of ELISA reading using phospho-PKA substrates antibody against peptides have Arginine or Lysine at -3 position.

Phospho-PKA Substrate Antibody

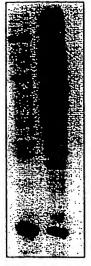
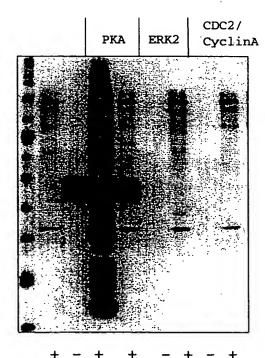


Figure 9: Western analysis of calyculin A-treated A431 cells using Phospho-PKA Substrate Antibody.

- + calyculin A



Pigure 10: Western analysis of A431 cell extracts phosphory-lated by protein kinase A, ERK2 or CDC2/cyclinA in vitro using Phospho-PKA Substrate Antibody.

Cell Extracts
PKI

Independent
Antibodies Using Peptide Libraries as Antigens
Comb, et al
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Phospho-Serine/Threonine Phenylalanine Antibody

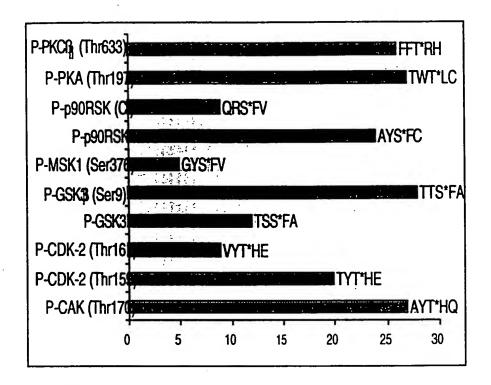


Figure 11: Signal to noise ratio of ELISA reading using phospho-Serine/threonine phenylalanine antibody aganist the peptides srounded by phenylalanine, tyrosine or tryptophan.

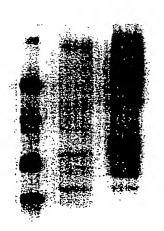


Figure 12: Western analysis of calyculin A-treated A431 cells using phospho-Serine/phenylalanine substates antibody.

Production of Motif-Specific Context-Independent Antibodies Using Peptide Libraries as Antigen Comb, et al Atty. Docket No: CST-138 CIP2 Sheet 16 of 23

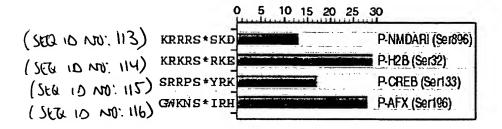


Figure 13. Signal to noise ratio of ELISA reading, using a context-independent antibody specific for the phospho-PKC consensus substrate motif, with phospho-PKC substrate containing peptides and nonphospho peptides.

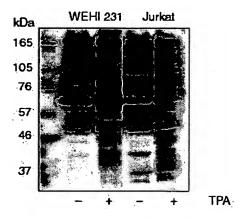


Figure 14. Western blot analysis of WEHI 231 cells or Jurkat cells untreated (–) or treated (+) with TPA, probed with a context-independent antibody specific for the phospho-PKC consensus substrate motif.

Production of Motif-Specific Context-Independent Antibodies Using Peptide Libraries as Antigens Comb, et al Atty. Docket No: CST-138 CIP2 Sheet 17 of 23

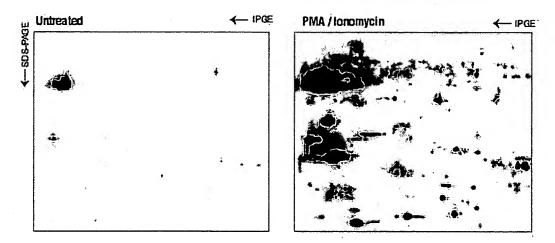


Figure 15. Western blot analysis of whole cell lysates of Jurkat cells untreated and treated with PMA (50 ng/ml) and ionomycin (1 μ M) for 20 minutes prior to lysis, using a context-independent antibody specific for the phospho-PKC consensus substrate motif. Proteins were separated by 2D electrophoresis prior to blotting.

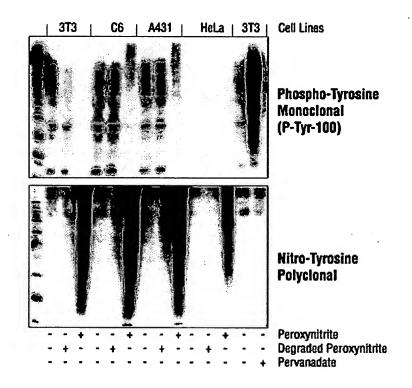


Figure 16. Western analysis of whole cell lysates of different cells untreated or treated with peroxynitrite, degraded peroxynitrite or pervanadate using a context-independent antibodies specific for phosphotyrosine (upper), and a polyclonal context-independent antibody specific for nitrotyrosine (lower).

Production of Motif-Specific Context-Independent Antibodies Using Peptide Libraries as Antigens Comb, et al Atty. Docket No: CST-138 CIP2 Sheet 18 of 23

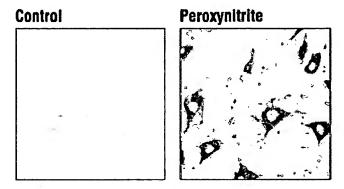


Figure 17. Immunocytochemical staining of NIH/3T3 cells treated with degraded peroxynitrite (control) or peroxynitrite using a polyclonal context-independent antibody specific for nitrotyrosine (brown).

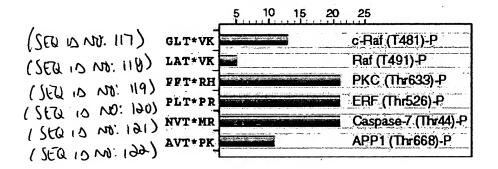


Figure 18. Phosphothreonine-X-arginine motif-specific context-independent antibody ELISAs: Signal to noise ratio of phospho versus nonphospho peptides containing the phospho-threonine-X-arginine motif. (T* denotes phosphorylated threonine.)

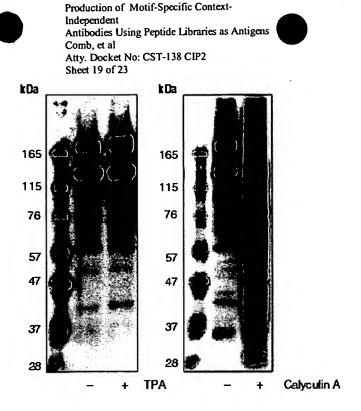


Figure 19. Western blot analysis of Jurkat cell extracts untreated (–) and treated (+) with TPA or Calyculin A, using a context-independent antibody specific for the phosphothreonine-X-arginine motif.

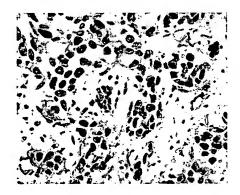


Figure 20. Immunohistochemical staining of proteins containing phosphorylated threonine-X-arginine motifs in paraffin-embedded human breast carcinoma, using a context-independent antibody specific for the motif.

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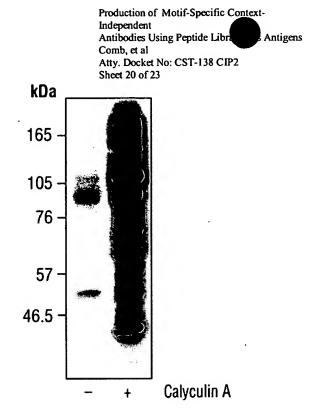


Figure 21. Western blot analysis of calyculin A treated A431 cells, using a context-independent antibody specific for the phospho-14-3-3 binding motif #2 (phospho(Ser)-Arg-X-(Tyr/Phe)-X-pSer).

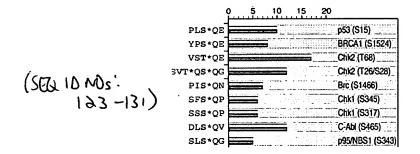
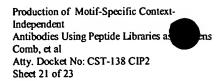


Figure 22. Phospho-ATM/ATR consensus substrate motif-specific, context-independent antibody ELISAs: Signal to noise ratio of phospho versus nonphospho peptides. (S* or T* denote phosphorylated serine or threonine.)



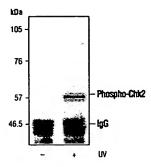


Figure 23. Chk2 transfected and UV treated COS cell extracts immunoprecipitated with Chk2 antibody then detected by Western blotting, using a context-independent antibody specific for phospho-ATM/ATR consensus substrate motif.

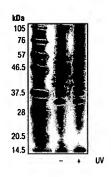


Figure 24. Western blot analysis of UV treated COS cells, using a context-independent antibody specific for phospho-ATM/ATR consensus substrate motif.

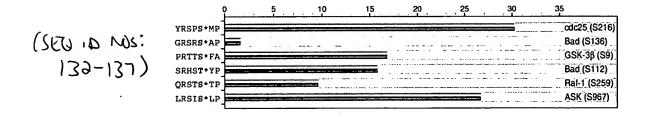


Figure 25. Phospho-14-3-3 binding motif-specific, context-independent monoclonal antibody ELISAs: Signal to noise ratio of phospho versus nonphospho 14-3-3 binding motif peptides. (T* and S* denote phosphorylated threonine and serine).

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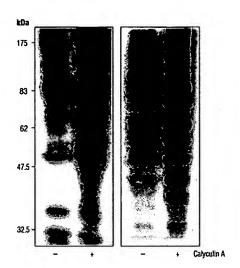


Figure 26. Western blot analysis of calyculin A treated A431 cells, using a context-independent antibodies specific for phospho-14-3-3 binding motif #1(left, monoclonal; right, polyclonal).

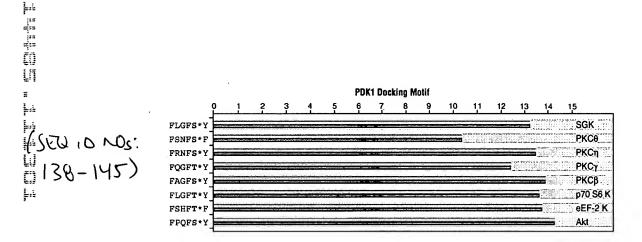


Figure 27. Phospho-PDK1 docking motif-specific, context-independent monoclonal antibody ELISAs: Signal to noise ratio of phospho versus non-phospho peptides corresponding to potential PDK1 docking motifs. (T* and S* denote phosphorylated threonine and serine.)

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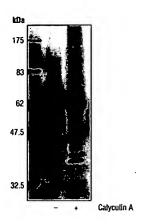


Figure 28. Western blot analysis of extracts from A431 cells untreated or treated with 0.1 μ M calyculin A for 30 minutes prior to lysis, using a monoclonal context-independent antibody specific for the phospho-PDK1 docking motif.

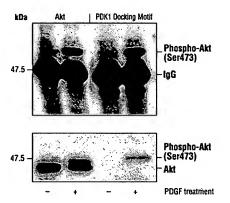


Figure 29. Immunoprecipitation of extracts from NIH/3T3 cells untreated or treated with 100 ng/ml of PDGF for 20 minutes prior to lysis, using a monoclonal context-independent antibody specific for phospho-PDK1 docking motif and an antibody specific for Akt, then probed with the PDK1 docking motif monoclonal antibody (upper) and the Akt antibody (lower).